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(54) **Unsaturated alcohols as accelerators for hydrosilation**

Ungesättigte Alkohole als Beschleuniger für Hydrosilylierung

Alcools insaturés comme accélérateurs pour l'hydrosilylation

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(56) References cited:

EP-A- 0 460 589 **EP-A- 0 510 957**
EP-A- 0 738 731 **EP-A- 0 751 140**
US-A- 2 823 218

- **CHEMICAL ABSTRACTS, vol. 110, no. 21, 22 May 1989 Columbus, Ohio, US; abstract no. 192835k, KORNETKA, Z.W.: "preparation of allyl 3-(4-morpholinyl)propionates" XP002035249 & PL 131 983 A (KORNETKA, Z.W.)**

Remarks:

The file contains technical information submitted after the application was filed and not included in this specification

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Description

[0001] The present invention is a hydrosilation process where a silicon hydride is contacted with an unsaturated reactant in the presence of a platinum catalyst and with an accelerator selected from unsaturated secondary and tertiary alcohols or sited unsaturated secondary and tertiary alcohols. The accelerators are especially useful for the hydrosilation of unsaturated reactants where the unsaturation is in the internal portion of the reactant's structure, for example, as in cyclopentene and cyclohexene. The accelerators are believed to be effective in the presence or absence of oxygen.

[0002] It is known in the art to produce organosilicon compounds by reacting a silicon hydride containing compound with an unsaturated organic compound in the presence of a catalyst. This reaction is typically referred to as hydrosilation or hydrosilylation. Typically the catalyst is platinum metal on a support, a platinum compound generally in a solvent or a platinum complex.

[0003] Representative of this prior art are US-As 2,823,218 and 3,220,972.

[0004] EP-A 0 460 589 discloses a method for the preparation of cyclopentyl trichlorosilane by the hydrosilation reaction of trichlorosilane and cyclopentene in the presence of a chlorine-deficient chloroplatinic acid catalyst. Suitable solvents for the catalyst include lower aliphatic alcohols, aromatic hydrocarbon solvents and silicone fluids.

[0005] One of the major problems with hydrosilation reactions is the deactivation of catalyst prior to completion of the reaction. One method for catalyst reactivation has been to expose the reaction mixture to oxygen. For example, US-A 4,578,497 discloses the use of an oxygenated platinum containing catalyst for use in hydrosilating alkylsilanes. Similarly, US-A 5,359,111 describes a method for controlling hydrosilation reaction mixtures by controlling the solution concentration of oxygen in the reaction mixture, relative to the concentration of platinum present.

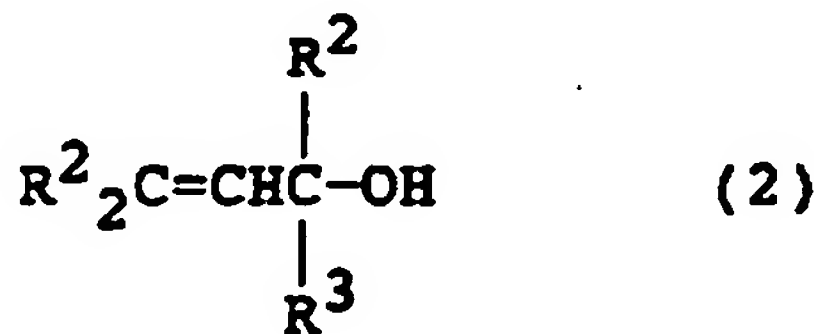
[0006] In addition to deactivation, hydrosilation processes of the art are not particularly effective in hydrosilating internal unsaturated bonds in organic molecules. We have unexpectedly found that unsaturated secondary and tertiary alcohols or sited unsaturated secondary and tertiary alcohols will act as accelerators for platinum catalyzed hydrosilation processes. These accelerators improve yield of the process in the presence, or absence, of oxygen and are particularly effective in facilitating the hydrosilation of internal unsaturated bonds of organic molecules.

[0007] Also, US-A 5,424,470 claims the use of unsaturated ketones as accelerators for hydrosilation reactions.

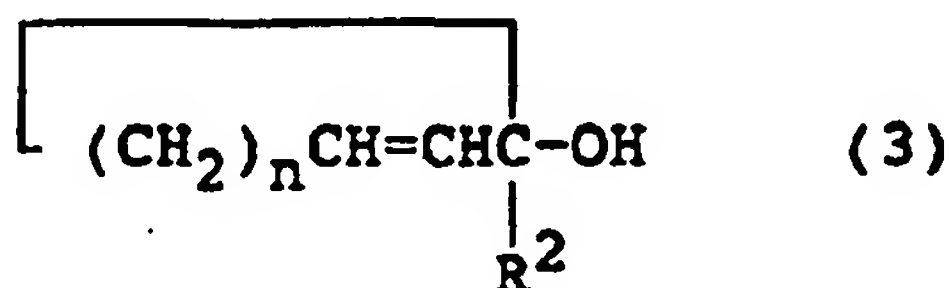
[0008] The present invention is a hydrosilation process where a silicon hydride is contacted with an unsaturated reactant in the presence of a platinum catalyst and with an accelerator. The hydrosilation process comprises contacting

(A) a silicon hydride described by formula $R^1_a H_b SiX_{4-a-b}$ (1) where each R^1 is independently selected from a group consisting of alkyls comprising one to 20 carbon atoms, cycloalkyls comprising four to 12 carbon atoms and aryls; each X is independently selected from halogen atom or organooxy radicals described by formula $-OR^1$, where R^1 is as previously described, $a=0$ to 3, $b=1$ to 3 and $a+b=1$ to 4; and

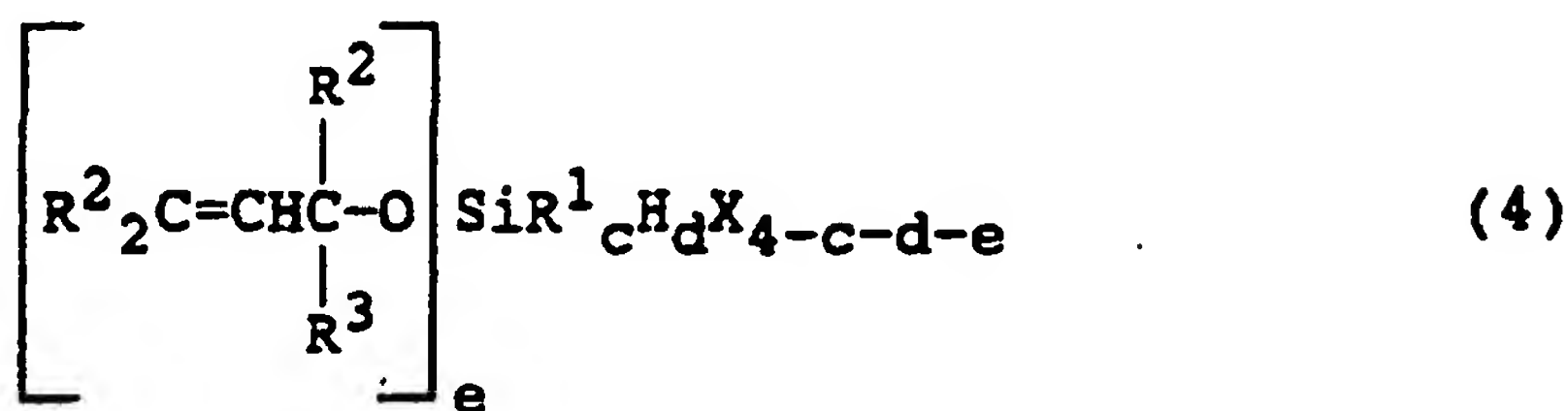
(B) an unsaturated reactant selected from a group consisting of (i) substituted and unsubstituted unsaturated organic compounds, (ii) silicon compounds comprising substituted or unsubstituted unsaturated organic substituents, and (iii) mixtures of (i) and (ii); in the presence of a platinum catalyst selected from platinum compounds or platinum complexes and an accelerator selected from unsaturated secondary and tertiary alcohols described by formulas



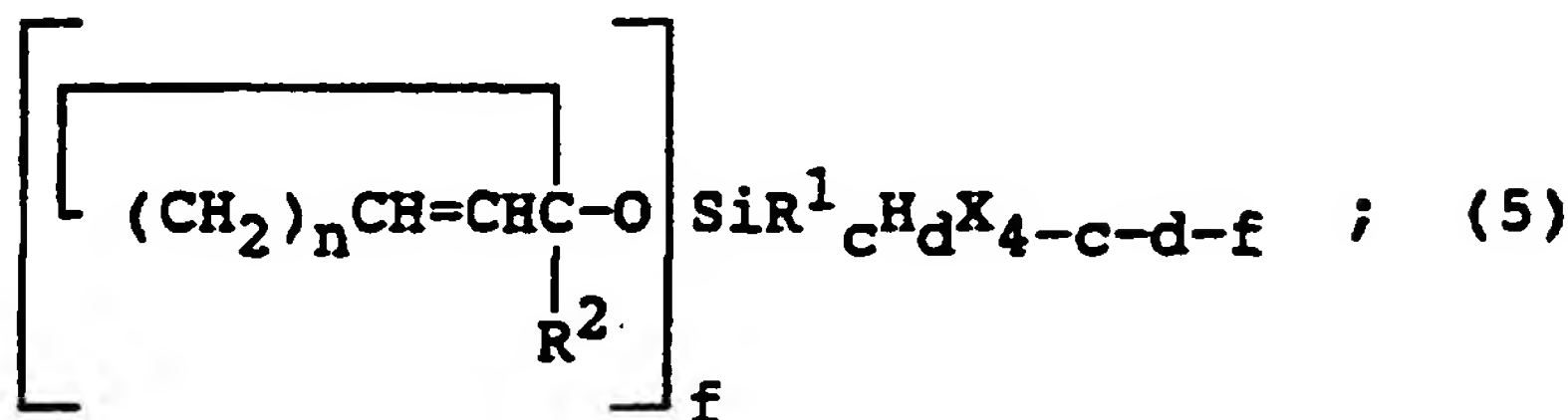
and



and silylated unsaturated secondary and tertiary alcohols described by formulas



and



where each R^2 is independently selected from hydrogen atom or alkyls comprising one to six carbon atoms, each R^3 is an independently selected alkyl comprising one to six carbon atoms, $c=0$ to 3, $d=0$ to 3, $c+d=0$ to 3, $e=1$ to 4, $f=1$ to 4 and $n=0$ to 10.

[0009] The contacting of the silicon hydride with the unsaturated reactant is effected in standard type reactors for conducting hydrosilation processes. The contact of reactants and subsequent reaction may be run as a continuous, semi-continuous or batch reaction.

[0010] Silicon hydrides which are useful in our process are described by formula (1), where each R^1 is independently selected from a group consisting of alkyls comprising one to 20 carbon atoms; cycloalkyls comprising four to 12 carbon atoms and aryls; $a=0$ to 3, $b=1$ to 3 and $a+b=1$ to 4.

[0011] In formula (1) it is preferred that each R^1 be independently selected from a group consisting of alkyls comprising one to six carbon atoms. Even more preferred is when each R^1 is methyl.

[0012] In formula 1, each X is independently selected from halogen atom or organooxy radicals described by formula $-\text{OR}^1$, where R^1 is as previously described. Preferred is when X is chlorine.

[0013] Examples, of silicon hydrides described by formula (1) which are useful in the present process include trimethylsilane, dimethylsilane, triethylsilane, dichlorosilane, trichlorosilane, methyldichlorosilane, dimethylchlorosilane, ethyldichlorosilane, cyclopentylchlorosilane, methylphenylchlorosilane, (3,3,3-trifluoropropyl)dichlorosilane and methylmethoxychlorosilane. A preferred silicon hydride of formula (1) is selected from methyldichlorosilane or dichlorosilane.

[0014] The silicon hydride is contacted with an unsaturated reactant selected from a group consisting of (i) substituted and unsubstituted unsaturated organic compounds, (ii) silicon compounds comprising substituted or unsubstituted unsaturated organic substituents and (iii) mixtures of (i) and (ii). For this invention, "unsaturated" means that the compound contains at least one carbon-carbon double bond.

[0015] Specific examples of the unsaturated reactants useful in this process include unsubstituted cycloalkene compounds comprising at least four carbon atoms, substituted cycloalkene compounds comprising at least four carbon atoms, linear alkene compounds comprising two to 30 carbon atoms, branched alkene compounds comprising four to 30 carbon atoms and mixtures of two or more of any of the above compounds.

[0016] The substituted and unsubstituted cycloalkene compounds useful in the present process are those containing one or more unsaturated carbon-carbon bonds in the ring. The unsubstituted cycloalkene compounds are, for example, cyclobutene, cyclopentene, cyclohexene, cycloheptene, cyclooctene, cyclopentadiene, 1,3-cyclohexadiene and 1,3,5-cycloheptatriene. Substituted unsaturated compounds include, for example, 3-methylcyclopentene, 3-chlorocyclobutene, 4-phenylcyclohexene and 3-methylcyclopentadiene. The preferred cycloalkene compounds are cyclohexene and cyclopentene, with cyclohexene being the most preferred.

[0017] Other unsaturated organic compounds useful in this process are linear and branched alkenyl compounds including, for example, compounds with terminal unsaturation such as 1-hexene and 1,5-hexadiene, compounds with

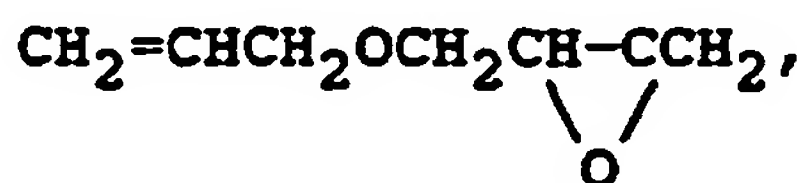
internal unsaturation such as trans-2-hexene and unsaturated aryl containing compounds such as styrene or α -methylstyrene.

[0018] The unsaturated reactants may also comprise halogen, oxygen in the form of acids, anhydrides, alcohols, esters and ethers; and nitrogen. Two or more of the above described unsaturated organic compounds may also be used in the present process.

[0019] The unsaturated organic compounds comprising halogen include, for example, vinyl chloride, allyl chloride, allyl bromide, allyl iodide, allyl bromide, methallyl chloride, trichloroethylene, tetrachloroethylene, tetrafluoroethylene, chloroprene, vinylidene chloride and dichlorostyrene.

[0020] Suitable unsaturated organic compounds comprising oxygen include, for example, ethers such as allyl and vinyl ethers; alcohols such as allyl alcohol (vinyl carbinol), methylvinylcarbinol and ethynyldimethylcarbinol; acids such as acrylic, methacrylic, vinylacetic, oleic, sorbic and linolenic; and esters such as vinyl acetate, allyl acetate, butenyl acetate, allyl stearate, methylacrylate, ethyl-crotonate, diallyl succinate and diallyl phthalate. Suitable nitrogen containing unsaturated organic compounds include, for example, indigo, indole, acrylonitrile and allyl cyanide.

[0021] Specifically included within the definition of unsaturated organic compounds are those substituted by organofunctional moieties such as $\text{CH}_2=\text{CHCH}_2\text{OC}(\text{O})\text{C}(\text{CH}_3)=\text{CH}_2$, $\text{CH}_2=\text{CHCH}_2\text{NHCH}_2\text{CH}_2\text{NH}_2$, $\text{CH}_2=\text{CHCH}_2\text{NH}_2$,



$\text{CH}_2=\text{CHCH}_2\text{SH}$, $\text{CH}_2=\text{CHSi}\{\text{O}(\text{CH}_2)_2\text{OCH}_3\}_3$ and $\text{CH}_2=\text{CHCH}_2\text{N}(\text{HCl})\text{HCH}_2\text{CH}_2\text{NHCH}_2(\text{C}_6\text{H}_4)\text{CH}=\text{CH}_2$.

[0022] The unsaturated organic compound is a silicon compound comprising substituted and unsubstituted organic substituents as described by, for example, formulas $(\text{CH}_2=\text{CH}(\text{CH}_2)_g)_h\text{R}^1\text{Si}(\text{OR}^1)_{4-h-i}$ and $(\text{CH}_2=\text{CH}(\text{CH}_2)_g)_h\text{R}^1\text{SiCl}_{4-h-i}$, where R^1 is as previously described, $g=0$ to 12 , $h=1$ to 3 , $i=0$ to 3 and $h+i=1$ to 4 .

[0023] Prior to contact of the silicon hydride with the unsaturated reactant, it may be preferable to treat or purify the unsaturated reactant. Useful methods are those known in the art for treating or purifying unsaturated organic compounds and include distillation and treatment with an adsorbent such as activated alumina or molecular sieves.

[0024] The relative amounts of silicon hydride and unsaturated reactant used in our process can be varied within wide limits. Although one unsaturated carbon-carbon linkage per silicon-bonded hydrogen atom is stoichiometric, there is no requirement that the process be run under stoichiometric conditions. Generally, it is preferred that the process

be run with a stoichiometric excess of silicon hydride. Preferred is when the process is run with 0.1 to ten percent stoichiometric excess of silicon hydride. However, for safety reasons, it may be preferred to run the process with an excess of unsaturated reactant, for example when the silicon hydride is dichlorosilane.

[0025] The silicon hydride and unsaturated reactant are contacted in the presence of a platinum catalyst selected from platinum compounds or platinum complexes. Any platinum containing material which effects the reaction between the silicon hydride and an unsaturated carbon-carbon bond of the unsaturated organic compound is useful in this process. Examples of useful platinum catalysts are described in US-As 4,578,497; 3,220,972 and 2,823,218.

[0026] The platinum catalyst can be, for example, chloroplatinic acid, chloroplatinic acid hexahydrate, Karstedt's catalyst (i.e., a complex of chloroplatinic acid with sym-divinyldimethylsiloxane), dichlorobis(triphenylphosphine) platinum(II), cis-dichlorobis(acetonitrile)platinum(II), dicarbonyldichloroplatinum(II), platinum chloride and platinum oxide.

[0027] A preferred platinum catalyst is selected from the group consisting of chloroplatinic acid, chloroplatinic acid hexahydrate and platinum vinylsiloxane complexes such as a neutralized complex of chloroplatinic acid or platinum dichloride with sym-divinyldimethylsiloxane.

[0028] Generally, those concentrations of platinum catalyst which provide at least one mole of platinum per billion moles of unsaturated carbon-carbon bonds, added to the process by the unsaturated reactant, is useful in the present process. Concentrations of platinum catalyst providing as high as one mole of platinum per one thousand moles of unsaturated carbon-carbon bonds added by the unsaturated reactant may be useful. Higher concentrations of platinum may be used if desired. A preferred concentration of platinum catalyst is that providing one to 1000 moles of platinum per 1×10^6 moles of unsaturated carbon-carbon bonds.

[0029] The platinum catalyst may be dissolved in a solvent for ease of handling and to facilitate measuring the small amounts typically needed. Suitable solvents include, for example, non-polar hydrocarbon solvents such as benzene, toluene and xylene or polar solvents such as alcohols, ketones, glycols and esters.

[0030] The present process is carried out in the presence of an accelerator selected from a group consisting of unsaturated secondary and tertiary alcohols, described by formulas (2) and (3) and silylated unsaturated secondary and

tertiary alcohols, described by formulas (4) and (5). In these formulae, each substituent R^2 is independently selected from hydrogen atom or alkyls comprising one to six carbon atoms. In addition to hydrogen, R^2 can be methyl, ethyl, propyl, butyl and hexyl. Preferred is when R^2 is selected from hydrogen atom or methyl. In formulae (2) and (4), each R^3 is an independently selected alkyl comprising one to six carbon atoms as described for R^2 . Preferred is when R^3 is methyl. In formulae (4) and (5), it is preferred that R^1 be methyl.

[0031] In formulae (3) and (5), n has a value of zero to 10. Preferred is when n is three. In the silylated unsaturated secondary and tertiary alcohols described by formulae (4) and (5), c has a value of zero to three; d has a value of zero to three; and c plus d has a value of zero to three. In formula (4), e has a value of one to four. Preferred is when e has a value of one. In formula (5), f has a value of one to 4. Preferred is when f has a value of one.

[0032] In formulae (4) and (5), each X is independently selected from halogen atom or organooxy radicals described by formula $-OR^1$, where R^1 is as previously described. Preferred is when X is a chlorine atom.

[0033] A preferred accelerator for use in the present process is selected from 2-methyl-3-buten-2-ol or silylated 2-methyl-3-buten-2-ol (i.e., 2-trimethylsiloxy-2-methyl-3-butene).

[0034] An effective concentration of the accelerator is added to the present process, where an effective concentration is one that facilitates initiation of the reaction between the silicon hydride and the unsaturated organic compound, accelerates the rate of the reaction or reduces loss of reactivity of the catalyst in the process. A useful effective concentration of the accelerator is generally within a range of 0.01 to 20 weight percent of the weight of the unsaturated reactant. Preferred is when the accelerator is 0.1 to ten weight percent of the weight of the unsaturated reactant. The accelerator is added to the process as a pre-mix with the platinum catalyst or is added separately.

[0035] The temperature at which the present process is conducted is generally within a range of -10°C . to 220°C . It is preferred to conduct our process at a temperature of 15°C . to 170°C . The most preferred temperature for the process is within a range of 30°C . to 150°C .

Example 1

[0036] A variety of alcohols were evaluated for their ability to accelerate the reaction of methyldichlorosilane with cyclohexene in the presence of a platinum catalyst.

[0037] A stock mixture was prepared in an argon purged and blanketed bottle. The stock mixture comprised four molar percent excess of methyldichlorosilane in cyclohexene which had been treated with 13X molecular sieves. Then, 3.7×10^{-5} to 6×10^{-5} moles of platinum, as a platinum divinylsiloxane complex, per mole of cyclohexene was added to the stock mixture. Aliquots of this catalyzed stock solution were then transferred to argon-purged glass tubes which contained alcohols as described in Table 1 at a concentration of one weight percent of alcohol per total cyclohexene added to the tube. The tubes were heat sealed under argon, purged and heated at 80°C . for three hours. Thereafter, the tubes were cooled and the contents analyzed by gas chromatography using a thermal conductivity detector (GC-TC). The results of this analysis are reported in Table 1 as the normalized area percent of methyl(cyclohexyl)dichlorosilane ($\text{MeC}_6\text{H}_{11}\text{SiCl}_2$) under the GC-TC trace minus the area of the cyclohexene as 100 percent. The results are given as the mean value for the number of runs indicated in parenthesis.

TABLE 1

Alcohols as Accelerators For Platinum Catalyzed Addition of MeHSiCl_2 to Cyclohexene	
Type Alcohol	Area% $\text{Me}(\text{C}_6\text{H}_{11})\text{SiCl}_2$
None	37.6 (7)
2-Methyl-3-buten-2-ol	93.5 (1)
Silylated 2-methyl-3-buten-2-ol	58.4 (1)
Allyl	5.9 (2)
Trans-2-hexen-1-ol	6.3 (4)
Hexyl	5.5 (2)

Example 2

[0038] The ability of unsaturated alcohols to accelerate the reaction of dichlorosilane with cyclopentene in the presence of a platinum catalyst was evaluated.

[0039] A stock mixture comprising 14.2 weight percent of dichlorosilane in cyclopentene was prepared in an argon purged and blanketed bottle. Aliquots of the mixture were transferred to argon purged glass tubes containing a platinum divinylsiloxane complex providing 7×10^{-4} moles of platinum per mole of dichlorosilane. Unsaturated alcohol or silylated unsaturated alcohol, providing two weight percent based on the total mass was then added to the tubes. The tubes

were heat sealed under an argon blanket and heated at 120°C. for one hour. Thereafter, the tubes were cooled and the contents analyzed by GC-TC. The results of the analysis are reported in Table 2 as the normalized area percent of cyclopentyldichlorosilane (CpHSiCl₂) and dicyclopentyldichlorosilane (Cp₂SiCl₂) under the GC-TC trace minus the area of the cyclopentene as 100 percent.

TABLE 2

Unsaturated Alcohols as Accelerators For Platinum Catalyzed Addition of Dichlorosilane to Cyclopentene		
Type Alcohol	Area% CpHSiCl ₂	Area% Cp ₂ SiCl ₂
None	56.9	0.0
2-Methyl-3-buten-2-ol	52.8	5.3
2-Trimethylsiloxy-2-methyl-3-butene	81.9	0.0

Example 3

[0040] The ability of 2-cyclohexen-1-ol to accelerate the reaction of methyldichlorosilane with cyclohexene in the presence of a platinum catalyst was evaluated. To an argon purged flask fitted with a mechanical stirrer and a condenser were added 78 g of cyclohexene which had been treated with 13X molecular sieves, 0.8 g of 2-cyclohexen-1-ol and a platinum divinylsiloxane complex providing 1.4×10^{-4} moles of platinum per mole of methyldichlorosilane. The mixture was heated under an argon purge to 80°C. and then 104 g of methyldichlorosilane were added at a rate of 2 ml per minute. During the addition of the methyldichlorosilane, an exothermic reaction occurred raising the temperature of the reaction mixture to 98°C. After cooling, the content of the flask was analyzed by GC-TC and found to contain 91.2 area percent of cyclohexylmethyldichlorosilane.

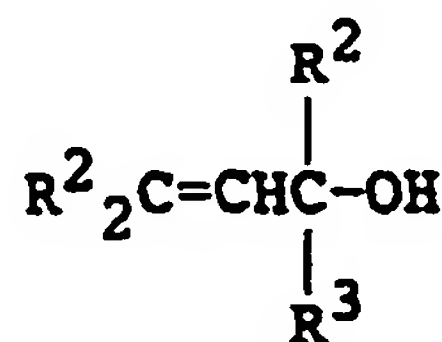
Claims**1. A hydrosilation process comprising contacting**

(A) a silicon hydride described by formula $R^1_a H_b SiX_{4-a-b}$ where each R^1 is independently selected from a group consisting of alkyls comprising one to 20 carbon atoms, cycloalkyls comprising four to 12 carbon atoms and aryls; each X is independently selected from halogen atom or organooxy radicals described by formula $-OR^1$, where R^1 is as previously described, $a=0$ to 3, $b=1$ to 3 and $a+b=1$ to 4; and

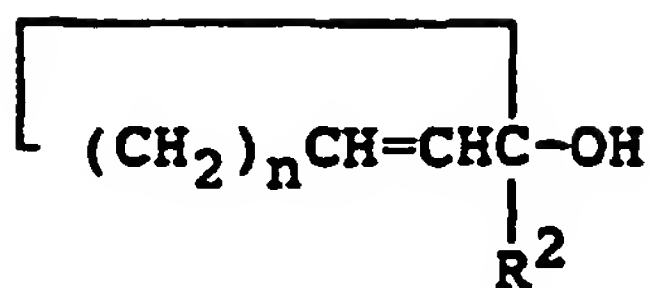
(B) an unsaturated reactant selected from a group consisting of

- (i) substituted and unsubstituted unsaturated organic compounds,
- (ii) silicon compounds comprising substituted or unsubstituted unsaturated organic substituents, and
- (iii) mixtures of (i) and (ii);

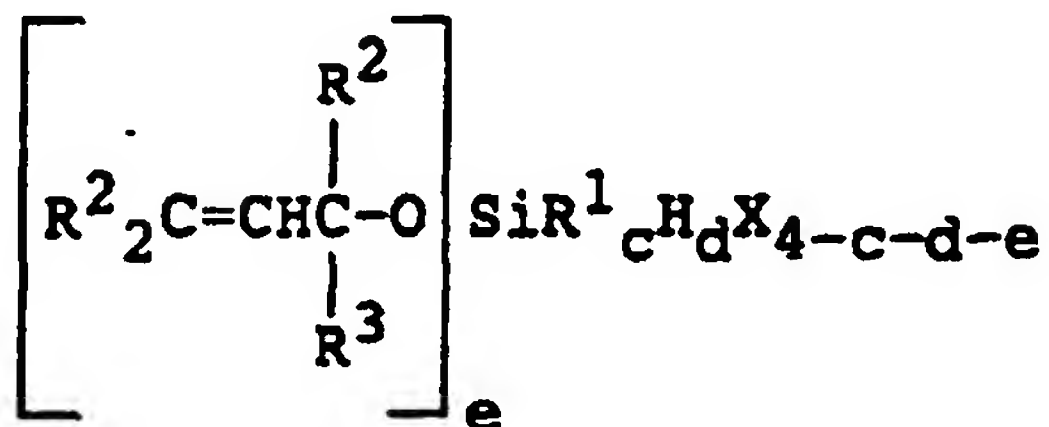
in the presence of a platinum catalyst selected from platinum compounds or platinum complexes and an accelerator selected from unsaturated secondary and tertiary alcohols described by formulas



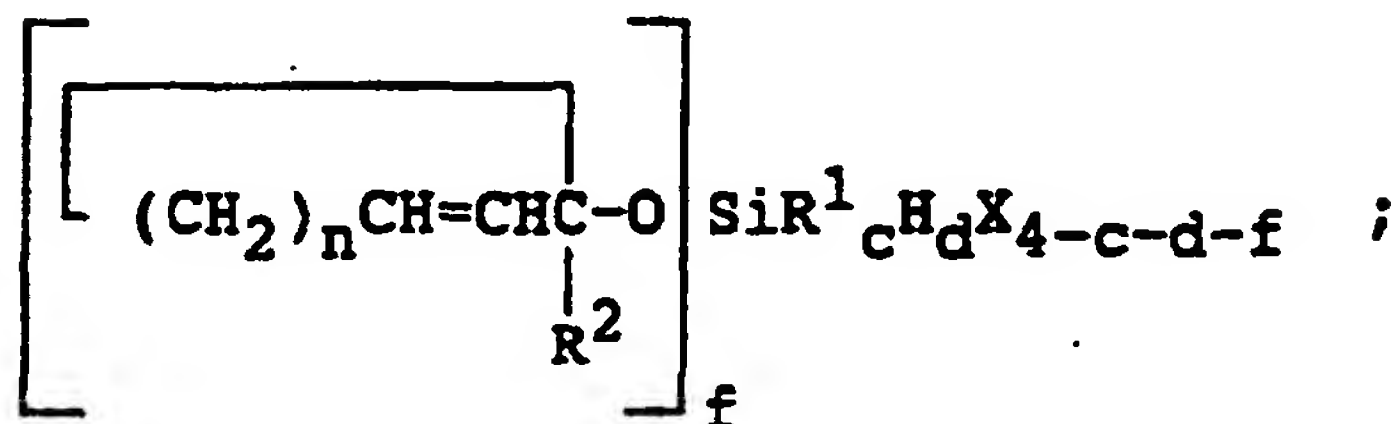
and



and silylated unsaturated secondary and tertiary alcohols described by formulas



and



where each R^2 is independently selected from hydrogen atom or alkyls comprising one to six carbon atoms, each R^3 is an independently selected alkyl comprising one to six carbon atoms, $c=0$ to 3, $d=0$ to 3, $c+d=0$ to 3, $e=1$ to 4, $f=1$ to 4 and $n=0$ to 10.

2. A process of claim 1 where 0.1 to ten percent stoichiometric excess of silicon hydride is contacted with the unsaturated reactant.
3. The process of claim 1 where the platinum catalyst provides to the process one to 1000 moles of platinum per 1×10^6 moles of unsaturated carbon-carbon bonds provided by the unsaturated reactant.
4. The process of claim 1 where the concentration of the accelerator is within a range of 0.01 to 20 weight percent of the weight of the unsaturated reactant.
5. The process of claim 1 where the silicon hydride is contacted with the unsaturated reactant at a temperature within a range of -10°C. to 220°C.

Patentansprüche

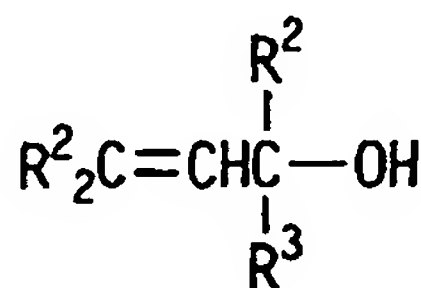
1. Hydrosilylierungsverfahren, umfassend Inberührungbringen von

(A) einem Siliciumhydrid, beschrieben durch Formel $\text{R}^1_a\text{H}_b\text{SiX}_{4-a-b}$, worin jedes R^1 unabhängig voneinander ausgewählt ist aus einer Gruppe, die aus Alkylen mit 1-20 Kohlenstoffatomen, Cycloalkylgruppen mit 4-12 Kohlenstoffatomen und Arylgruppen besteht; jedes X unabhängig voneinander ausgewählt ist aus einem Halogenatom oder Organooxyresten, beschrieben durch Formel $-\text{OR}^1$, worin R^1 wie zuvor definiert ist, a gleich 0 bis 3 ist, b gleich 1 bis 3 ist und $a+b$ gleich 1 bis 4 ist; und

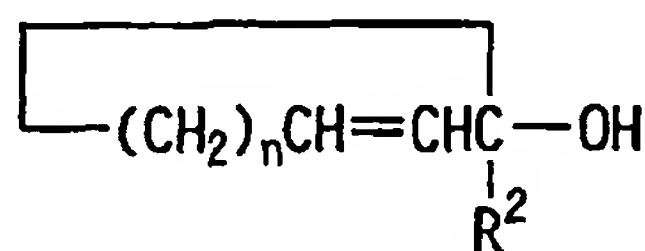
(B) einem ungesättigten Reaktanten, ausgewählt aus einer Gruppe bestehend aus

- (i) substituierten und unsubstituierten ungesättigten organischen Verbindungen,
- (ii) Siliciumverbindungen, die substituierte oder unsubstituierte ungesättigte organische Substituenten enthalten, und
- (iii) Mischungen aus (i) und (ii);

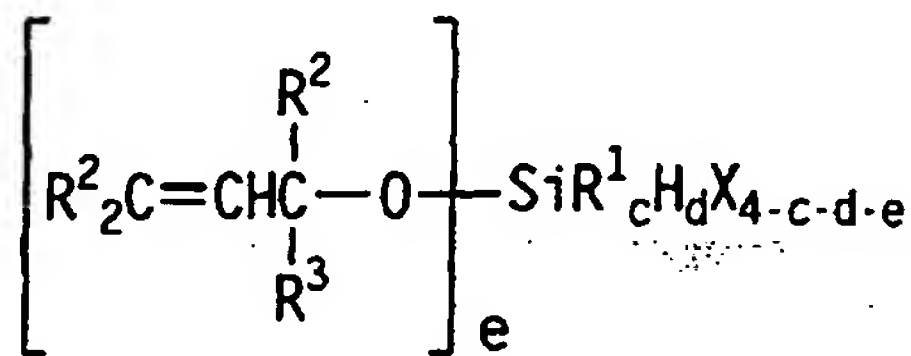
in Gegenwart eines Platinkatalysators, ausgewählt aus Platinverbindungen oder Platinkomplexen, und einem Beschleuniger, ausgewählt aus ungesättigten sekundären und tertiären Alkoholen, beschrieben durch Formeln:



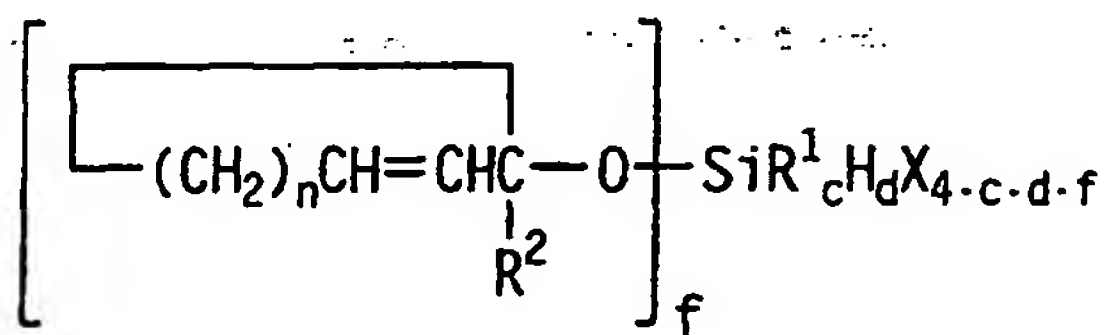
und



und silylierten ungesättigten sekundären und tertiären Alkoholen, beschrieben durch Formeln:



und



worin jedes R^2 unabhängig voneinander ausgewählt ist aus einem Wasserstoffatom oder Alkylgruppen mit 1-6 Kohlenstoffatomen, jedes R^3 ein unabhängig voneinander ausgewähltes Alkyl mit 1-6 Kohlenstoffatomen ist, c gleich 0 bis 3 ist, d gleich 0 bis 3 ist, c+d gleich 0 bis 3 ist, e gleich 1 bis 4 ist, f gleich 1 bis 4 ist und n gleich 0 bis 10 ist.

2. Verfahren nach Anspruch 1, worin 0,1-10% stöchiometrischer Überschuss von Siliciumhydrid mit dem ungesättigten Reaktanten in Berührung gebracht werden.
3. Verfahren nach Anspruch 1, worin der Platinkatalysator dem Prozess 1-1000 mol Platin pro 1×10^6 mol ungesättigter Kohlenstoff-Kohlenstoff-Bindungen, die von dem ungesättigten Reaktanten geliefert werden, zur Verfügung stellt.
4. Verfahren nach Anspruch 1, wobei die Konzentration des Beschleunigers innerhalb eines Bereichs von 0,01-20 Gew.-% des Gewichts des ungesättigten Reaktanten liegt.

5. Verfahren nach Anspruch 1, worin das Siliciumhydrid mit dem ungesättigten Reaktanten bei einer Temperatur innerhalb eines Bereichs von -10°C bis 220°C in Berührung gebracht wird.

5 Revendications

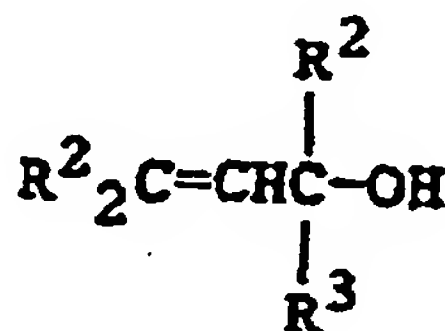
1. Procédé d'hydrosilation comprenant la mise en contact de

(A) un hydrure de silicium décrit par la formule $R^1_a H_b SiX_{4-a-b}$ dans laquelle chaque R^1 est choisi indépendamment dans un groupe formé par les alkyle comprenant de 1 à 20 atomes de carbone, les cycloalkyle comprenant de 4 à 12 atomes de carbone et des aryle ; chaque X étant choisi indépendamment parmi un atome d'halogène ou les radicaux organoxy décrits par la formule $-OR^1$, dans laquelle R^1 est comme précédemment décrit, $a = 0$ à 3, $b = 1$ à 3 et $a+b = 1$ à 4 ; et

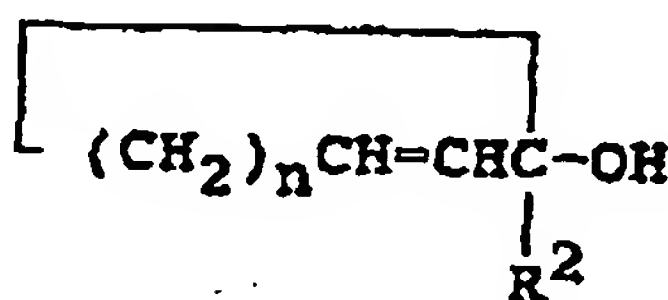
(B) un réactif insaturé choisi dans le groupe formé par :

- (i) des composés organiques insaturés substitués et non substitués,
- (ii) des composés du silicium comprenant des substituants organiques insaturés substitués ou non substitués, et
- (iii) des mélanges de (i) et de (ii) ;

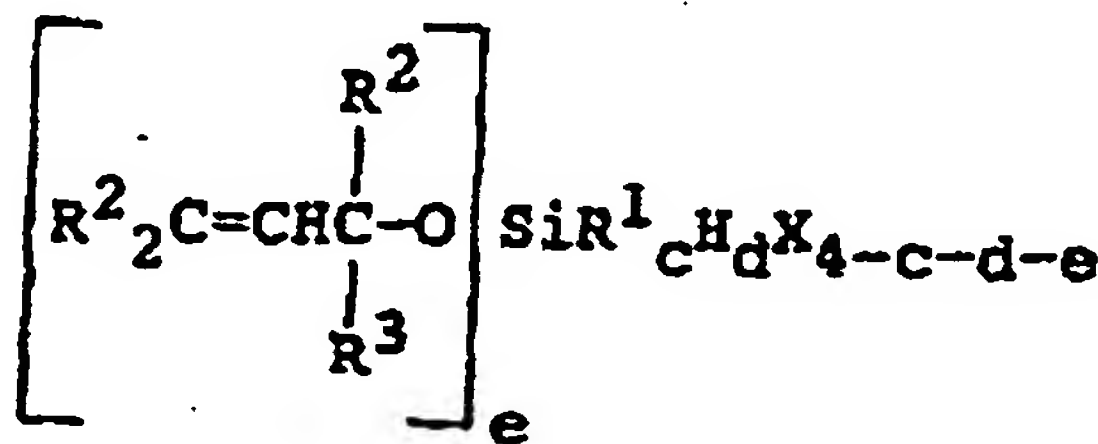
en présence d'un catalyseur à base de platine choisi parmi des composés du platine ou des complexes de platine et un accélérateur choisi parmi des alcools secondaires et tertiaires insaturés, décrits par les formules :



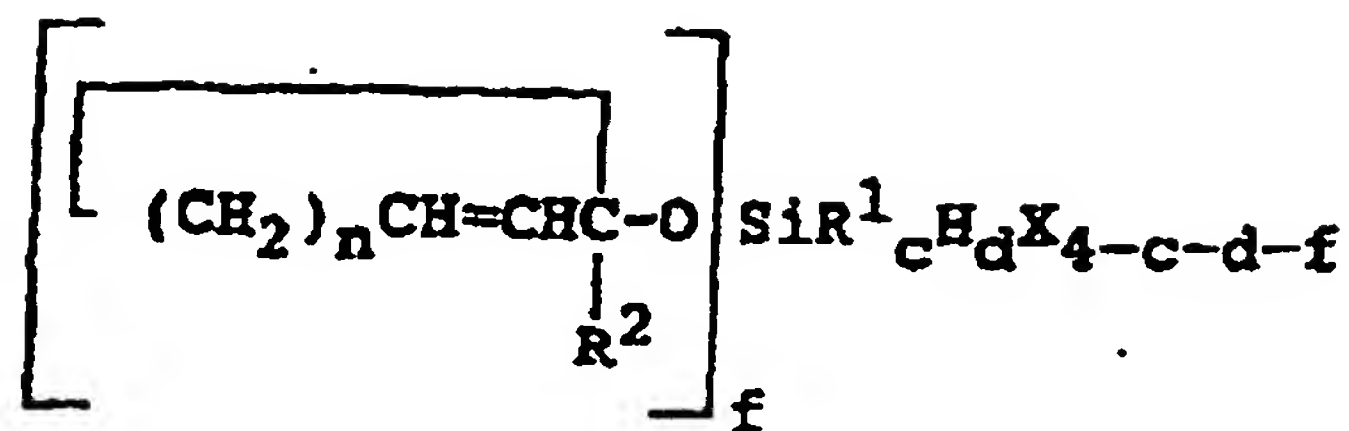
et



et des alcools secondaires et tertiaires insaturés silatés décrits par les formules :



et



dans lesquelles chaque R^2 est choisi indépendamment parmi un atome d'hydrogène ou des alkyle comprenant de 1 à 6 atomes de carbone, chaque R^3 indépendamment est un alkyle comprenant 1 à 6 atomes de carbone, $c = 0$ à 3, $d = 0$ à 3, $c+d = 0$ à 3, $e = 1$ à 4, $f = 1$ à 4 et $n = 0$ à 10.

2. Procédé selon la revendication 1, caractérisé en ce qu'un excès stoechiométrique de 0,1 à 10 pour cent d'hydruure de silicium est mis en contact avec le réactif insaturé.
3. Procédé selon la revendication 1, caractérisé en ce que le catalyseur à base de platine fournit au procédé une à 1000 moles de platine pour 1×10^6 moles de liaisons carbone-carbone insaturées apportées par le réactif insaturé.
4. Procédé selon la revendication 1, caractérisé en ce que la concentration de l'accélérateur est comprise entre 0,01 et 20 pour cent en poids par rapport au poids du réactif insaturé.
5. Procédé selon la revendication 1, caractérisé en ce que l'hydruure de silicium est mis en contact avec le réactif insaturé à une température comprise entre -10°C et 220°C .

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